

(19)



JAPANESE PATENT OFFICE

PATENT ABSTRACTS OF JAPAN

(11) Publication number: 2000212180 A

(43) Date of publication of application: 02.08.00

(51) Int. Cl

**C07D401/12**  
**A61P 1/04**  
**A61P 31/04**  
**A61K 31/4709**  
**A61K 31/4725**  
**A61K 31/5377**  
**C07D401/14**  
**C07D413/14**

(21) Application number: 11012632

(22) Date of filing: 21.01.99

(71) Applicant: **WELFIDE CORP**

(72) Inventor: **SANO MITSUHARU**  
**YOKOYAMA YOSHIHITO**  
**KITANI HIROYUKI**  
**SAKURAI NOBUHIRO**  
**EBARA HIDEJI**  
**MIYOSHI MIE**

**(54) QUINOLINE COMPOUND**

**(57) Abstract:**

**PROBLEM TO BE SOLVED:** To obtain a new compound expressing a selective antibacterial activity against bacteria of the genus *Helicobacter* represented by *Helicobacter pylori* and useful for preventing and curing diseases such as gastritis, stomach ulcer and the likes caused by *Helicobacter* bacteria.

**SOLUTION:** This quinoline compound is a compound expressed by formula I {R1 is H, a halogen, nitro, cyano, an alkyl, an alkoxy, OH or the likes; R2 is H, a (halo)alkyl, a hydroxylalkyl or

the likes; R3 and R4 are each H, a halogen or an alkyl; X is O, S, SO or the likes; B is O, S, SO2 or the likes; D is D1-E [D1 is a single bond, a (substituted) alkylene or the likes; E is a (halo) alkoxyalkyl or the likes] or the likes}, e.g. 4-[(3-methyl-4-(2-morpholinoethylthio)-2-pyridyl)methoxy]quinoline 3 hydrochloride. The compound of formula I is produced by a method, etc., for reacting a compound of formula II with a compound (an acid adduct) of formula III [W is a reaction active atom or group (e.g. a halogen)].

**COPYRIGHT:** (C)2000,JPO

BEST AVAILABLE COPY

BEST AVAILABLE COPY

